

## EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
S75 7	1572	thiosemicarbazone	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:21
S75 8	76168	vaccine	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:21
S75 9	132321	adjuvant	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:21
S76 0	2	S757 same S758 same S759	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:24
S76 1	14	paul.in. near3 barsanti.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:24
S76 2	11	nathan.in. near3 brammeier.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:25
S76 3	3	anthony.in. near3 diebes.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:26
S76 4	21	liana.in. near3 lagniton.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:26
S76 5	28	simon.in. near3 ng.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:27
S76 6	32	Zhi-Jie.in. near3 Ni.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:28
S76 7	3	Casey.in. near3 Philbin.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:29

## EAST Search History

S76 8	21	Nicholas.in. near3 Valiante.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:32
S76 9	28	Allan.in. near3 Wagman.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:33
S77 0	29	Weibo.in. near3 Wang.in.	US-PGPUB; USPAT; EPO; JPO; DERWENT	AND	ON	2007/05/15 14:33

## STICSearch\_SEQID2.txt

## RESULT 3

AAB14903

ID AAB14903 standard; peptide; 24 AA.

XX

AC AAB14903;

XX

DT 08-JAN-2001 (first entry)

XX

DE Ht31 peptide.

XX

KW Calcineurin-binding peptide; A-kinase anchor protein; AKAP; AKAP 79;  
 KW immunostimulant; interleukin 2 expression modulation; graft rejection;  
 KW transplantation; T cell-mediated disorder.

XX

OS Unidentified.

XX

PN US6107104-A.

XX

PD 22-AUG-2000.

XX

PF 27-SEP-1996; 96US-00721458.

XX

PR 23-NOV-1994; 94US-00344227.

PR

PR 15-MAR-1995; 95US-00404731.

PR

PR 17-JUL-1995; 95US-00503226.

XX

PA (ICOS-) ICOS CORP.

XX

PI Lockerbie RO, Gallatin WM, Lai Y, Howard ML;

XX

DR WPI; 2000-578541/54.

XX

PT Novel calcineurin deletion mutant having calcineurin polypeptide sequence

PT and binding A-kinase anchor proteins, for treating graft rejection

PT following organ transplantation and T cell-mediated disorders.

XX

PS Example 9; Col 5-6; 53pp; English.

XX

CC The present sequence was used in an example to demonstrate that  
 CC association of cAMP-dependent protein kinase (PKA) with an anchoring  
 CC protein in T cells modulates IL-2 production. A-kinase anchoring protein  
 CC 79 (AKAP 79) binds both PKA and calcineurin and so co-localises a kinase  
 CC and a phosphatase that may regulate flux through a specific signalling  
 CC pathway. Calcineurin is a Ca<sup>2+</sup>/calmodulin-dependent protein phosphatase  
 CC which is involved in many intracellular signalling pathways. It  
 CC participates in regulation of IL-2 expression following T cell  
 CC stimulation in T cells. Calcineurin-binding peptides derived from AKAP 79  
 CC may be used to inhibit calcineurin activity in a cell. The peptides are  
 CC useful for treating graft rejection following organ transplantation and  
 CC for treating T cell-mediated disorders. Calcineurin deletion mutants  
 CC which bind AKAP 79 are useful for defining an AKAP 79 binding site, for  
 CC stimulating the immune response, stimulating activated T cells for  
 CC selected clonal expansion, or for enhancing T cell responses to  
 CC experimental stimuli for evaluation of early events in T cell biology and  
 CC activation of the immune response

XX

SQ Sequence 24 AA;

Query Match 100.0%; Score 111; DB 3; Length 24;

Best Local Similarity 100.0%; Pred. No. 2.1e-10;

Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DLIEEAASRIVDAVIEQVKAAGAY 24

|||||

Db 1 DLIEEAASRIVDAVIEQVKAAGAY 24

STICSearchStrategy.txt

=> d que 152

```
L2      869 SEA FILE=REGISTRY ABB=ON PLU=ON (100708-37-0/BI OR
101091-29-6/BI OR 103907-05-7/BI OR 108-49-6/BI OR
109-09-1/BI OR 165047-24-5/BI OR 1722-12-9/BI OR 17532-19-3
/BI OR 17754-90-4/BI OR 22078-59-7/BI OR 22725-58-2/BI OR
23945-44-0/BI OR 2417-72-3/BI OR 2929-81-9/BI OR 301176-96-
5/BI OR 301176-97-6/BI OR 301202-34-6/BI OR 301348-19-6/BI
OR 301348-20-9/BI OR 301348-21-0/BI OR 301349-64-4/BI OR
301349-65-5/BI OR 301350-41-4/BI OR 301351-29-1/BI OR
302563-23-1/BI OR 302564-39-2/BI OR 302564-43-8/BI OR
302564-45-0/BI OR 303058-86-8/BI OR 303759-55-9/BI OR
303782-53-8/BI OR 309277-46-1/BI OR 316130-78-6/BI OR
316136-61-5/BI OR 326901-08-0/BI OR 33342-19-7/BI OR
33342-21-1/BI OR 340220-71-5/BI OR 340226-07-5/BI OR
340299-81-2/BI OR 340309-70-8/BI OR 340316-53-2/BI OR
343591-25-3/BI OR 344944-35-0/BI OR 345616-23-1/BI OR
34721-90-9/BI OR 34721-92-1/BI OR 34721-94-3/BI OR
347335-07-3/BI OR 348594-85-4/BI OR 354535-04-9/BI OR
355414-88-9/BI OR 3608-75-1/BI OR 3608-81-9/BI OR 371941-52
-5/BI OR 3731-53-1/BI OR 375361-02-7/BI OR 38594-22-8/BI
OR 403657-28-3/BI OR 413571-56-9/BI OR 418783-85-4/BI OR
420816-07-5/BI OR 443144-51-2/BI OR 464207-10-1/BI OR
468749-49-7/BI OR 51940-64-8/BI OR 56621-48-8/BI OR
60698-40-0/BI OR 64248-62-0/BI OR 6839-88-9/BI OR 6868-27-5
/BI OR 6868-28-6/BI OR 69045-79-0/BI OR 712349-33-2/BI OR
725726-27-2/BI OR 725726-28-3/BI OR 725726-29-4/BI OR
725726-30-7/BI OR 725726-31-8/BI OR 725726-32-9/BI OR
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725726-42-1/BI OR 725726-43-2/BI OR 725726-44-3/BI OR
725726-45-4/BI OR 725726-46-5/BI OR 725726-47-6/BI OR
725726-48-7/BI OR 725726-49-8/BI OR 725726-50-1/BI OR
725726-51-2/BI OR 725726-52-3/BI OR 725726-53-4/BI OR
725726-
L7      244 SEA FILE=REGISTRY ABB=ON PLU=ON "C7 H8 N4 S"/MF
L8      1 SEA FILE=REGISTRY ABB=ON PLU=ON L2 AND L7
L11     STR
```

NODE ATTRIBUTES:  
 DEFAULT MLEVEL IS ATOM  
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:  
 RING(S) ARE ISOLATED OR EMBEDDED  
 NUMBER OF NODES IS 8

STEREO ATTRIBUTES: NONE  
 L13 6896 SEA FILE=REGISTRY SSS FUL L11  
 L16 189 SEA FILE=HCAPLUS ABB=ON PLU=ON L8  
 L17 17 SEA FILE=HCAPLUS ABB=ON PLU=ON L16 AND THU/RL  
 L22 STR

VAR G1=X/9/11/N/AK  
 NODE ATTRIBUTES:  
 DEFAULT MLEVEL IS ATOM  
 DEFAULT ECLEVEL IS LIMITED

## STICSearchStrategy.txt

GRAPH ATTRIBUTES:  
RING(S) ARE ISOLATED OR EMBEDDED  
NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

L24 4889 SEA FILE=REGISTRY SUB=L13 SSS FUL L22  
L25 2905 SEA FILE=HCAPLUS ABB=ON PLU=ON L24  
L33 45 SEA FILE=HCAPLUS ABB=ON PLU=ON BARSANTI, P?/AU  
L34 6 SEA FILE=HCAPLUS ABB=ON PLU=ON BRAMMEIER, N?/AU  
L35 5 SEA FILE=HCAPLUS ABB=ON PLU=ON DIEBES, A?/AU  
L36 12 SEA FILE=HCAPLUS ABB=ON PLU=ON LAGNITON, L?/AU  
L37 2474 SEA FILE=HCAPLUS ABB=ON PLU=ON NG, S?/AU  
L38 881 SEA FILE=HCAPLUS ABB=ON PLU=ON NI, Z?/AU  
L39 257 SEA FILE=HCAPLUS ABB=ON PLU=ON PFISTER, K?/AU  
L40 17 SEA FILE=HCAPLUS ABB=ON PLU=ON PHILBIN, C?/AU  
L41 54 SEA FILE=HCAPLUS ABB=ON PLU=ON VALIANTE, N?/AU  
L42 49 SEA FILE=HCAPLUS ABB=ON PLU=ON WAGMAN, A?/AU  
L43 23333 SEA FILE=HCAPLUS ABB=ON PLU=ON WANG, W?/AU  
L44 456 SEA FILE=HCAPLUS ABB=ON PLU=ON WEINER, A?/AU  
L45 8 SEA FILE=HCAPLUS ABB=ON PLU=ON (L33 OR L34 OR L35 OR L36  
OR L37 OR L38 OR L39 OR L40 OR L41 OR L42 OR L43 OR L44)  
AND L25  
L46 16 SEA FILE=HCAPLUS ABB=ON PLU=ON L17 NOT L45  
L49 98 SEA FILE=HCAPLUS ABB=ON PLU=ON L16 AND PREP/RL  
L50 24 SEA FILE=HCAPLUS ABB=ON PLU=ON L49 AND PHARM?/SC,SX  
L51 23 SEA FILE=HCAPLUS ABB=ON PLU=ON L50 NOT L45  
L52 34 SEA FILE=HCAPLUS ABB=ON PLU=ON L51 OR L46

L52 ANSWER 34 OF 34 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1971:11774 HCAPLUS Full-text

DOCUMENT NUMBER: 74:11774

TITLE: Carcinostatic activity of thiosemicarbazones of  
formyl heteroaromatic compounds. VII.  
2-Formylpyridine derivatives bearing additional  
ring substituents

AUTHOR(S): French, Frederic A.; Blanz, Erwin J., Jr.;  
DoAmaral, Jefferson R.; French, Douglas A.

CORPORATE SOURCE: Chemother. Res. Lab., Mt. Zion Hosp. Med. Center,  
Palo Alto, CA, USA

SOURCE: Journal of Medicinal Chemistry (1970), 13(6),  
1124-30

CODEN: JMCMAR; ISSN: 0022-2623

DOCUMENT TYPE: Journal

LANGUAGE: English

ED Entered STN: 12 May 1984

GI For diagram(s), see printed CA Issue.

AB Sixteen 5-substituted-2-formylpyridine thiosemicarbazones were synthesized and tested

for antitumor activity against 5 mouse tumor systems, such as L-1210 leukemia and the Lewis  
lung carcinoma. The I, II, and III were extremely potent inhibitors of tumor-derived  
ribonucleotide diphosphate reductase and inhibited the synthesis of DNA. At 100 mg/kg/day  
i.p., I had a broad spectrum of antitumor activity in addition to a significant cure rate in  
L-1210 leukemia.

IT 3608-75-1  
(neoplasm inhibition by)

RN 3608-75-1 HCAPLUS

CN Hydrazinecarbothioamide, 2-(2-pyridinylmethylene)- (CA INDEX NAME)

DialogSearchStrategy.txt

? b 155,10,5,73,34

[File 155] MEDLINE(R) 1950-2007/May 11

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[File 10] AGRICOLA 70-2007/May

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[File 5] Biosis Previews(R) 1926-2007/May W1

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\*File 5: BIOSIS has been enhanced with archival data. Please see HELP NEWS 5 for information.

[File 73] EMBASE 1974-2007/May 10

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[File 34] SciSearch(R) Cited Ref Sci 1990-2007/Apr W4

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? d s

Set	Items	Description
S1	4663	S THIOSEMICARBAZONE
S2	391079	S VACCINE
S3	391079	S VACCINE
S4	241856	S ADJUVANT
S5	0	S S1 (S) S2 (S) S4
S6	0	S S1 AND S2 AND S4
S7	0	S S1 (S) S4
S8	4	S S1 AND S4
S9	4	RD (unique items)
S10	14	S "PYRIDINE-2-CARBALDEHYDE THIOSEMICARBAZONE"
S11	14	RD (unique items)
S12	0	S S4 AND S11